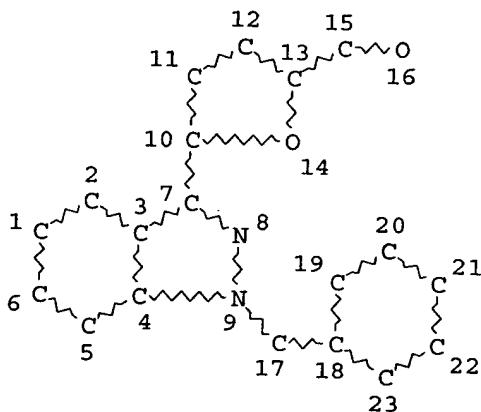


=> d que stat 123
 L1 STR



NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L3 139 SEA FILE=REGISTRY SSS FUL L1
 L21 1 SEA FILE=REGISTRY ABB=ON PLU=ON 170632-47-0
 L22 138 SEA FILE=REGISTRY ABB=ON PLU=ON L3 NOT L21
 L23 36 SEA FILE=HCAPLUS ABB=ON PLU=ON L22

=> d 123 ibib abs hitstr 1-36

L23 ANSWER 1 OF 36 HCAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2005:1341973 HCAPLUS

TITLE: The design and synthesis of YC-1 analogs as probes for soluble guanylate cyclase

AUTHOR(S): Hering, Kirk W.; Artz, Jennifer D.; Pearson, William H.; Marletta, Michael A.

CORPORATE SOURCE: Department of Chemistry, University of Michigan, Ann Arbor, MI, 48109-1055, USA

SOURCE: Bioorganic & Medicinal Chemistry Letters (2006), 16(3), 618-621

PUBLISHER: CODEN: BMCLE8; ISSN: 0960-894X
 Elsevier B.V.

DOCUMENT TYPE: Journal
 LANGUAGE: English

AB Soluble guanylate cyclase (sGC) is highly activated in the presence of both YC-1 [1-benzyl-3-(5-hydroxymethyl-2-furyl)indazole] and CO. The design, synthesis, and activity (i.e., sGC activation) of photolabile analogs of YC-1 are presented. Initial results with 6-azido-3-(5-hydroxymethyl-2-furyl)-1-benzylindazole led to the synthesis of a tritium-labeled analog. When photoactivated, this analog labeled the α -subunit of sGC.

IT 876365-94-5P

RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant)

=> d his ful

(FILE 'HOME' ENTERED AT 14:15:25 ON 13 MAR 2006)

FILE 'REGISTRY' ENTERED AT 14:16:41 ON 13 MAR 2006

L1 STR

L2 10 SEA SSS SAM L1

L3 139 SEA SSS FUL L1

FILE 'HCAPLUS' ENTERED AT 14:18:20 ON 13 MAR 2006

L4 170 SEA ABB=ON PLU=ON L3

FILE 'REGISTRY' ENTERED AT 14:19:11 ON 13 MAR 2006

L5 STR L1

L6 121 SEA SUB=L3 SSS FUL L5

FILE 'HCAPLUS' ENTERED AT 14:20:07 ON 13 MAR 2006

L7 170 SEA ABB=ON PLU=ON L6

FILE 'REGISTRY' ENTERED AT 14:20:12 ON 13 MAR 2006

L8 STR L5

L9 0 SEA SUB=L3 SSS SAM L8

L10 1 SEA SUB=L3 SSS FUL L8

D SCA

L11 STR L8

L12 93 SEA SUB=L3 SSS FUL L11

FILE 'HCAPLUS' ENTERED AT 14:23:32 ON 13 MAR 2006

L13 170 SEA ABB=ON PLU=ON L12

FILE 'BEILSTEIN' ENTERED AT 14:24:50 ON 13 MAR 2006

L14 0 SEA SSS SAM L8

L15 0 SEA SSS FUL L8

FILE 'MARPAT' ENTERED AT 14:25:13 ON 13 MAR 2006

L16 0 SEA SSS SAM L8

L17 2 SEA SSS FUL L8

FILE 'HCAPLUS' ENTERED AT 14:25:30 ON 13 MAR 2006

L18 1 SEA ABB=ON PLU=ON L10

FILE 'MARPAT' ENTERED AT 14:25:34 ON 13 MAR 2006

L19 1 SEA ABB=ON PLU=ON L17 NOT L18

FILE HOME

FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 12 MAR 2006 HIGHEST RN 876514-29-3

DICTIONARY FILE UPDATES: 12 MAR 2006 HIGHEST RN 876514-29-3

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* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

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FILE HCPLUS

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FILE COVERS 1907 - 13 Mar 2006 VOL 144 ISS 12
FILE LAST UPDATED: 12 Mar 2006 (20060312/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

FILE BEILSTEIN
FILE LAST UPDATED ON JANUARY 17, 2006

FILE COVERS 1771 TO 2005.
FILE CONTAINS 9,428,406 SUBSTANCES

>>>PLEASE NOTE: Reaction Data and substance data are stored in separate documents and can not be searched together in one query. Reaction data for BEILSTEIN compounds may be displayed immediately with the display codes PRE (preparations) and REA (reactions). A substance answer set retrieved after the search for a chemical name, a compounds with available reaction information by combining with PRE/FA, REA/FA or more generally with RX/FA. The BEILSTEIN Registry Number (BRN) is the link between a BEILSTEIN compound and belonging reactions. For more detailed reaction searches BRNs can be searched as reaction partner BRNs Reactant BRN (RX.RBRN) or Product BRN (RX.PBRN).<<<

>>> FOR SEARCHING PREPARATIONS SEE HELP PRE <<<

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* SET NOTICE FEATURE: THE COST ESTIMATES CALCULATED FOR SET NOTICE
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* PATENT NUMBERS (PN) AND BABS ACCESSION NUMBERS (BABSAN) CAN NOW BE
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* NEW DISPLAY FORMATS ALLREF, ALLP AND BABSAN SHOW ALL REFERENCES,
ALL PATENT REFERENCES, OR ALL BABS ACCESSION NUMBERS FOR A
COMPOUND AT A GLANCE.

FILE MARPAT

FILE CONTENT: 1910-PRESENT VOL 144 ISS 11 (20060310/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1910-1987

MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
(COVERAGE TO THESE DATES IS NOT COMPLETE):

US	2006030554	09	FEB	2006
DE	102004053311	05	JAN	2006
EP	1609846	28	DEC	2005
JP	2006003337	05	JAN	2006
WO	2006012333	02	FEB	2006
GB	2415429	28	DEC	2005
FR	2873371	27	JAN	2006
RU	2266908	27	DEC	2005
CA	2495134	23	DEC	2005

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FILE CONTENT: 1910-PRESENT VOL 144 ISS 11 (20060310/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1910-1987

MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
(COVERAGE TO THESE DATES IS NOT COMPLETE):

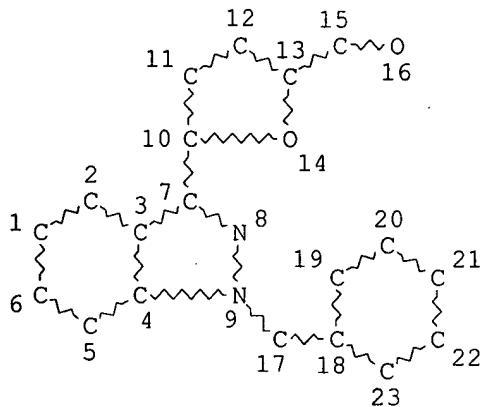
US	2006030554	09	FEB	2006
DE	102004053311	05	JAN	2006
EP	1609846	28	DEC	2005
JP	2006003337	05	JAN	2006
WO	2006012333	02	FEB	2006
GB	2415429	28	DEC	2005

FR 2873371 27 JAN 2006
 RU 2266908 27 DEC 2005
 CA 2495134 23 DEC 2005

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=> d que stat 118
 L1 STR



NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

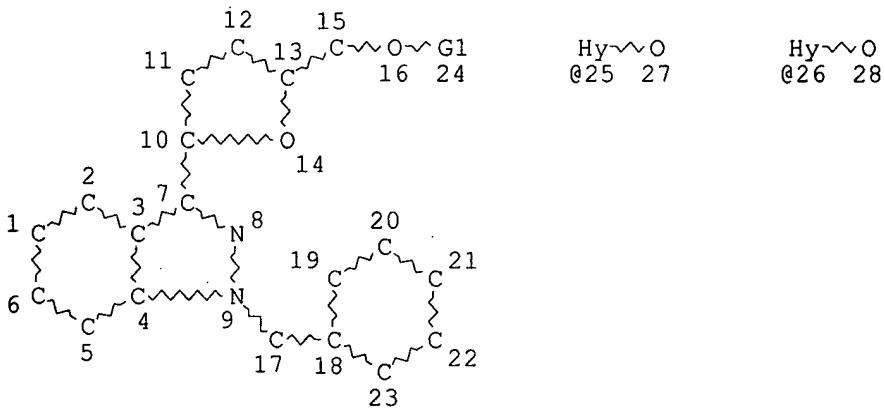
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L3 139 SEA FILE=REGISTRY SSS FUL L1
 L8 STR



VAR G1=25/26

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

GGCAT IS MCY SAT AT 25
 GGCAT IS MCY SAT AT 26
 DEFAULT ECLEVEL IS LIMITED
 ECOUNT IS E5 C E1 O AT 25
 ECOUNT IS E4 C E1 O AT 26

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 28

STEREO ATTRIBUTES: NONE

L10 1 SEA FILE=REGISTRY SUB=L3 SSS FUL L8
 L18 1 SEA FILE=HCAPLUS ABB=ON PLU=ON L10

=> d 118 ibib abs hitstr

YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L18 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:902206 HCAPLUS
 DOCUMENT NUMBER: 141:388641
 TITLE: HIF-1 expression inhibition-based method for
 inhibiting tumor angiogenesis and tumor growth
 INVENTOR(S): Park, Jong-Wan; Chun, Yang-Sook; Kim, Jinho
 PATENT ASSIGNEE(S): Bizbiotech Co., Ltd., USA
 SOURCE: PCT Int. Appl., 69 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004091648	A1	20041028	WO 2004-US10327	20040401
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004198798	A1	20041007	US 2003-407136	20030407
US 2005096370	A1	20050505	US 2003-642363	20030814
PRIORITY APPLN. INFO.:			US 2003-407136	A 20030407
			US 2003-642363	A 20030814

OTHER SOURCE(S): MARPAT 141:388641

AB The invention provides methods and pharmaceutical compns. for inhibiting expression of HIF-1 and HIF-1-regulated genes, angiogenesis, tumor growth, or tumor progression/metastasis, comprising contacting the tumor cells or tissue with a composition comprising 3-(5'-hydroxymethyl-2'-furyl)-1-benzylindazole, or a derivative thereof.

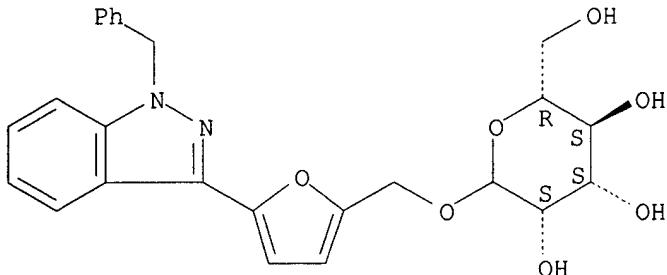
IT 781663-33-0

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (HIF-1 expression inhibition-based method for inhibiting tumor angiogenesis and tumor growth)

RN 781663-33-0 HCAPLUS

CN D-Mannopyranoside, [5-[1-(phenylmethyl)-1H-indazol-3-yl]-2-furanyl]methyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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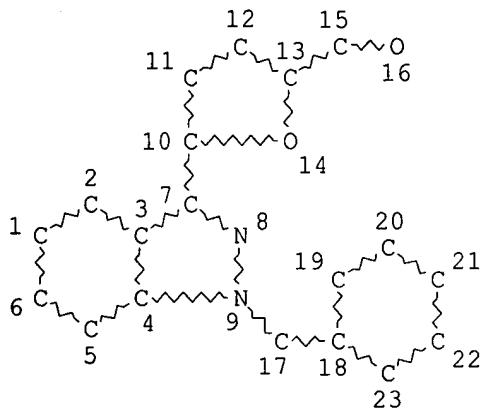
MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES (COVERAGE TO THESE DATES IS NOT COMPLETE):

US 2006030554 09 FEB 2006
 DE 102004053311 05 JAN 2006
 EP 1609846 28 DEC 2005
 JP 2006003337 05 JAN 2006
 WO 2006012333 02 FEB 2006
 GB 2415429 28 DEC 2005
 FR 2873371 27 JAN 2006
 RU 2266908 27 DEC 2005
 CA 2495134 23 DEC 2005

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=> d que stat 119
 L1 STR



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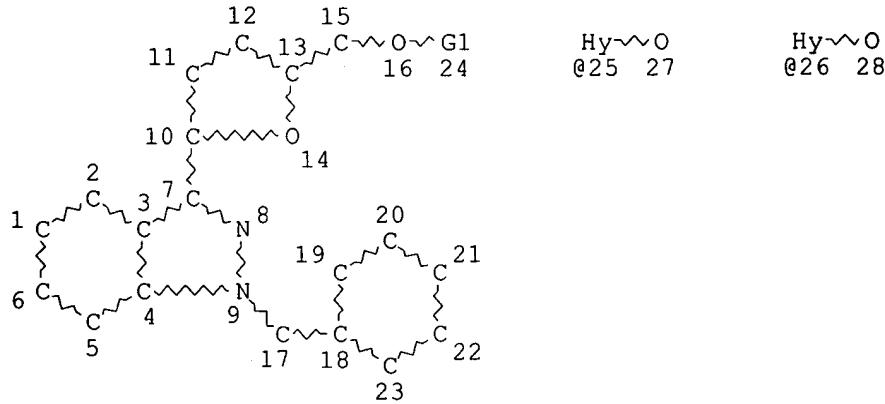
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 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L3 139 SEA FILE=REGISTRY SSS FUL L1
 L8 STR



VAR G1=25/26

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM
 GGCAT IS MCY SAT AT 25
 GGCAT IS MCY SAT AT 26
 DEFAULT ECLEVEL IS LIMITED
 ECOUNT IS E5 C E1 O AT 25
 ECOUNT IS E4 C E1 O AT 26

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 28

STEREO ATTRIBUTES: NONE

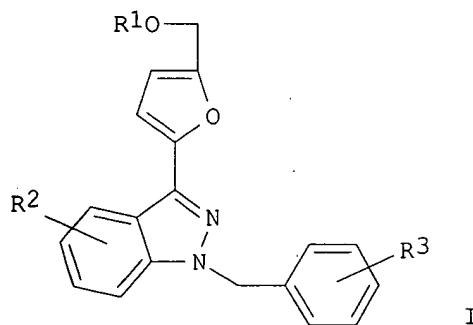
L10 1 SEA FILE=REGISTRY SUB=L3 SSS FUL L8
 L17 2 SEA FILE=MARPAT SSS FUL L8
 L18 1 SEA FILE=HCAPLUS ABB=ON PLU=ON L10
 L19 1 SEA FILE=MARPAT ABB=ON PLU=ON L17 NOT L18

=> d 119 ibib abs qhit

L19 ANSWER 1 OF 1 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 142:441845 MARPAT
 TITLE: Method for inhibiting tumor angiogenesis and tumor growth
 INVENTOR(S): Park, Jong-Wan; Chun, Yang-Sook; Kim, Jinho
 PATENT ASSIGNEE(S): Bizbiotech Co., Ltd., S. Korea
 SOURCE: U.S. Pat. Appl. Publ., 36 pp., Cont.-in-part of U.S. Ser. No. 407,136.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005096370	A1	20050505	US 2003-642363	20030814
US 2004198798	A1	20041007	US 2003-407136	20030407
WO 2004091648	A1	20041028	WO 2004-US10327	20040401
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			US 2003-407136	20030407
			US 2003-642363	20030814

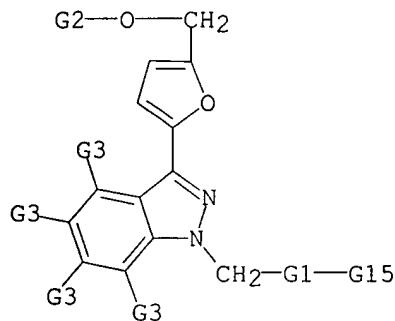
GI



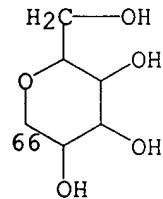
AB The present invention provides methods and pharmaceutical compns. for

inhibiting expressions of HIF-1 and HIF-1-regulated genes, angiogenesis, tumor growth, or tumor progression/metastasis comprising contacting the tumor cells or tissue with a composition comprising 3-(5'-hydroxymethyl-2'-furyl)-1-benzylindazole, or mixture of compds. of the formula I (where R1 = polyol, R2, R3 = H, alkyl, alkoxy, halogen, etc.).

MSTR 1



G1 = phenylene
 G2 = 66



Patent location: claim 7
 Note: substitution is restricted
 Note: additional ring and oxo formation also disclosed
 Note: and pharmaceutically acceptable solvates and salts
 Stereochemistry: and isomers and mixtures of isomers